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Visible-light-driven photoelectrochemical and photocatalytic performances of Cr-doped SrTiO₃/TiO₂ heterostructured nanotube arraysZhengbo Jiao¹, Tao Chen¹, Jinyan Xiong¹, Teng Wang¹, Gongxuan Lu¹, Jinhua Ye² & Yingpu Bi¹¹State Key Laboratory for Oxo Synthesis & Selective Oxidation, and National Engineering Research Center for Fine Petrochemical Intermediates, Lanzhou Institute of Chemical Physics, CAS, Lanzhou 730000, China, ²International Center for Materials Nanoarchitectonics (MANA), and Research Unit for Environmental Remediation Materials, National Institute for Materials Science, Tsukuba, 305-0047, Japan.

Well-aligned TiO₂ nanotube arrays have become of increasing significance because of their unique highly ordered array structure, high specific surface area, unidirectional charge transfer and transportation features. However, their poor visible light utilization as well as the high recombination rate of photoexcited electron-hole pairs greatly limited their practical applications. Herein, we demonstrate the fabrication of visible-light-responsive heterostructured Cr-doped SrTiO₃/TiO₂ nanotube arrays by a simple hydrothermal method, which facilitate efficient charge separation and thus improve the photoelectrochemical as well as photocatalytic performances.

One-dimensional TiO₂ array architectures, especially well-aligned TiO₂ nanotube arrays, exhibit promising photoelectric and photocatalytic performances in solar cells, water splitting and photocatalytic systems because of their unique highly ordered array structure, high specific surface area, unidirectional charge transfer and transportation features^{1–5}. However, their poor visible light utilization as well as the high recombination rate of photoexcited electron-hole pairs greatly limit their practical applications^{6–8}. Thereby, much effort has been devoted to the design and construction of novel hetero-composite structures to address these issues, such as metal or non-metal ion doping, noble metal loading, and hetero-coupling^{9–12}. In terms of semiconductor heterojunctions, SrTiO₃, whose conduction band edge is 200 mV more negative than TiO₂, have been considered as an alternate candidate for improving photoelectrochemical and photocatalytic performances by shifting the Fermi level of TiO₂ nanostructure to more negative potentials^{13–17}. Additionally, the coupling of SrTiO₃ and TiO₂ into a proper hetero-architecture can effectively facilitate the separation of photogenerated electrons and holes between SrTiO₃ and TiO₂, and their performances can be greatly enhanced¹⁸. However, note that another crucial issue about the visible light utilization still remains in these hetero-nanostructures as a result of their both relatively large band gap (3.2 eV)¹³. In view of efficient solar energy utilization, the design and fabrication of visible-light-responsive SrTiO₃/TiO₂ heterostructure nanotube arrays is highly desired because about 43% of solar energy is visible light while only about 4% is UV light¹⁹. Simultaneously, doping transition metals with SrTiO₃ material for narrowing its band gap and enhancing absorption of visible light have attracted considerable attention^{20–22}. In particular, chromium doping is of great interest because the occupied Cr 3d level is much lower (2.2 eV) than the conduction band bottom of SrTiO₃ determined by the Ti 3d orbital^{23,24}. However, the Cr-doped SrTiO₃/TiO₂ heterostructured nanotube arrays for facilitating the charge separation and enhancing the visible light response have rarely been reported up to now.

Herein, we demonstrate a facile synthetic strategy for fabricating uniform Cr-doped SrTiO₃/TiO₂ heterostructured nanotube arrays by in situ hydrothermal method using TiO₂ nanotube arrays as both template and precursor, and their structure and composition can be rationally tailored by simply adjusting the reaction time. Furthermore, it has been found that the Cr-doped SrTiO₃/TiO₂ heterostructure nanotube arrays exhibit much higher photoelectrochemical as well as photocatalytic performances than both pure TiO₂ nanotube arrays and SrTiO₃/TiO₂ nanotube arrays under visible light irradiation.

SUBJECT AREAS:
NANOSCALE MATERIALS
ENVIRONMENTAL CHEMISTRYReceived
15 July 2013Accepted
4 September 2013Published
23 September 2013Correspondence and
requests for materials
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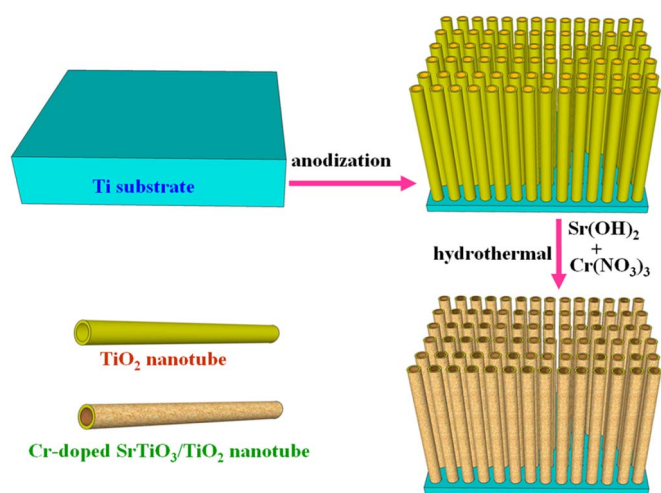


Figure 1 | Schematic illustration of the synthetic approach of Cr-doped SrTiO₃/TiO₂ heterostructure nanotube arrays.

Results

The fabrication approach of heterostructured Cr-doped SrTiO₃/TiO₂ nanotube arrays have been illustrated in Figure 1. Firstly, the well aligned TiO₂ nanotube arrays were produced by electrochemical anodization of Ti foil in hydrofluoric medium and then annealed at 500°C for 1 h in air to induce the transformation from amorphous to high crystalline anatase. Secondly, Cr-doped SrTiO₃/TiO₂ heterostructure nanotube arrays were synthesized by reacting TiO₂ nanotube arrays with Sr(OH)₂ in the presence of Cr(NO₃)₃ under hydrothermal treatment, in which TiO₂ nanotube arrays were utilized as both template and reactant.

Fig. 2A and Fig. 2B exhibit the typical scanning electron microscopy (SEM) images of the TiO₂ nanotube arrays prepared by electrochemical anodization and then anneal while keeping the tubular

architectures. It can be observed that TiO₂ nanotube arrays consist of uniform nanotubes with an average length of 3.8 μm, a mean pore diameter of 56 nm, and a wall thickness of about 35 nm (as shown in Fig. S1). The orifices and the side surfaces of these nanotubes are composed of relatively smooth surfaces without secondary nanostructures. However, when TiO₂ nanotube arrays were dipped into an aqueous solution that contains Sr(OH)₂ and Cr(NO₃)₃ under hydrothermal treatment, heterostructured Cr-doped SrTiO₃/TiO₂ nanotube arrays were obtained. Different from TiO₂ nanotube arrays surrounded by smooth surfaces, the side surfaces of Cr-doped SrTiO₃/TiO₂ heterostructure nanotube arrays become rough, and the orifices are cracked due to the penetration of Sr and Cr cations into TiO₂ nanotube arrays (as shown in Fig. 2C and 2D). Furthermore, the amount of Cr-doped SrTiO₃ can be rationally tailored by simply adjusting the reaction time (Fig. S2). It can be clearly seen from Fig. S2A that at the hydrothermal reaction time equal to 0.5 h, the side surfaces of the heterostructured Cr-doped SrTiO₃/TiO₂ nanotube arrays are only a little crude. However, further increasing the reaction time up to 1.5 or 2 h, the surfaces of as-fabricated Cr-doped SrTiO₃/TiO₂ heterostructures gradually become coarser, which can be ascribed to the excessive transformation of TiO₂ to Cr-doped SrTiO₃.

Additionally, Fig. 2E shows the transmission electron microscopy (TEM) image of the fragments of Cr-doped SrTiO₃/TiO₂ nanocomposite obtained through the sonication treatment in the presence of acetone solution. To further determine the specific distribution of Cr, Sr, Ti and O elements, high angle annular dark field scanning TEM (HAADF-STEM) measurement of the Cr-doped SrTiO₃/TiO₂ heterostructure nanotube arrays has been conducted (Fig. 2F). As shown in Fig. 2G–J, it can be clearly seen that the distribution of Cr, Sr and Ti elements is homogeneous and uniform, indicating the well dispersal of Cr-doped SrTiO₃ on the outer wall of TiO₂ nanotubes. Note that the distribution of O element exhibits a larger area than the other elements, which may be ascribed to the impurity from the copper grid. Besides, the amount distribution of each element has

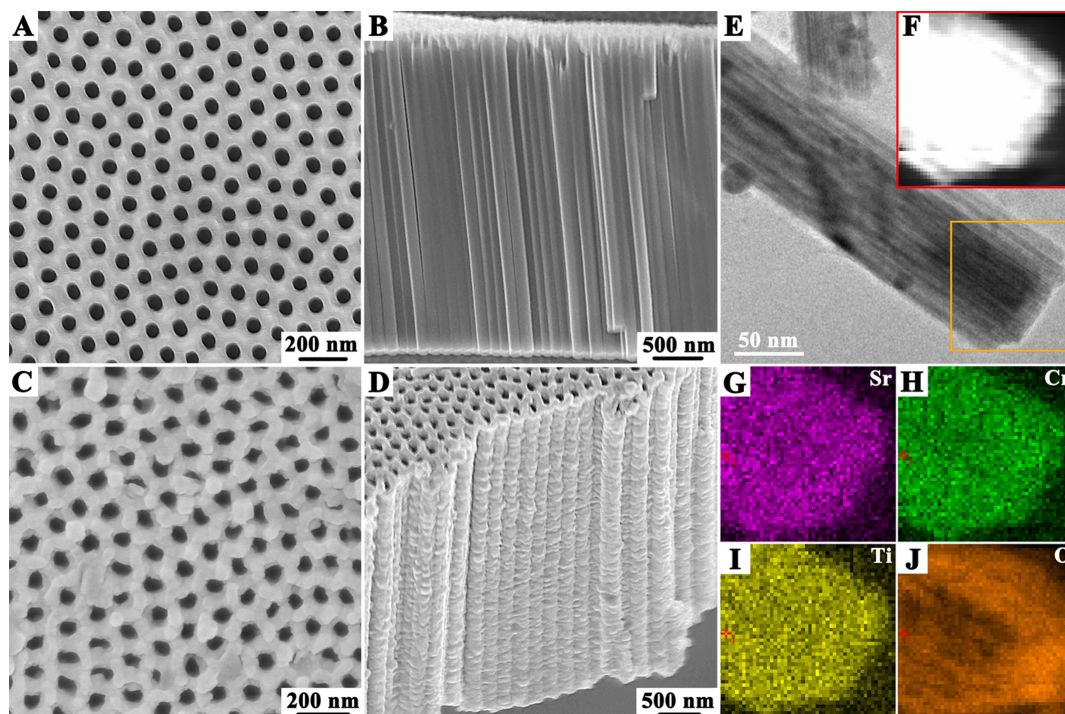


Figure 2 | SEM images (A, B) of TiO₂ nanotube arrays and (C, D) heterostructured Cr-doped SrTiO₃/TiO₂ nanotube arrays treated with 1 h hydrothermal reaction times. (E) TEM image, (F) HAADF-STEM image and (G–J) EDX elemental mapping images of the 1 h treated Cr-doped SrTiO₃/TiO₂ heterostructure.



been detected by the energy dispersive X-ray (EDX) spectrum and shown in Fig. S3.

The UV-Vis diffusive absorption spectra of as-prepared TiO_2 nanotube arrays and Cr-doped $\text{SrTiO}_3/\text{TiO}_2$ heterostructures with various growth stages have been shown in Fig. 3A. It can be observed that comparing with TiO_2 nanotube arrays, there are evident red-shifts in the absorbance peak edges of these heterostructures by increasing the hydrothermal reaction time, which should be due to the amount increase of Cr-doped SrTiO_3 . Their crystal structure and composition have been further studied by X-ray diffraction (XRD) and the results have been shown in Fig. 3B, which clearly reveals that the peak intensity of Cr-doped SrTiO_3 increases gradually with prolonging the hydrothermal times. Moreover, note that the XRD patterns of the Cr-doped SrTiO_3 are nearly identical with the cubic SrTiO_3 (Fig. S5), indicating that Cr cations doping did not introduce any impurities²⁴. The comparison of (110) diffraction peaks between $\text{SrTiO}_3/\text{TiO}_2$ and Cr-doped $\text{SrTiO}_3/\text{TiO}_2$ heterostructure nanotube arrays (Fig. S6) shows that the peak position of Cr-doped $\text{SrTiO}_3/\text{TiO}_2$ nanocomposites shifts slightly toward a higher degree, implying the partial substitution of Cr^{3+} cations for Sr^{2+} in the Cr-doped $\text{SrTiO}_3/\text{TiO}_2$ heterostructures²³, which may be due to the smaller ionic radius of Cr^{3+} (0.062 nm) than that of Sr^{2+} (0.118 nm) but a slightly larger than that of Ti^{4+} (0.061 nm). Consequently, it can be concluded that the right or left shift of the peak position generally represents the substitution of Cr^{3+} cations for Sr^{2+} or Ti^{4+} , respectively²³.

Furthermore, X-ray photoelectron spectroscopy (XPS) measurements were carried out to examine the valence states of chromium element in Cr-doped $\text{SrTiO}_3/\text{TiO}_2$ heterostructure nanotube arrays. As shown in Fig. 3C, the fully scanned spectrum confirms the existence of Cr, Ti, O, Sr and C elements. The high-resolution Cr 2p spectrum shown in Fig. 3D reveals that the doublet peaks at 576.9

and 586.5 eV could be indexed to $\text{Cr } 2p_{3/2}$ and $\text{Cr } 2p_{1/2}$, respectively. In the Cr 2p spectrum, the sharp peak at 576.9 eV is assigned to the trivalent chromium corresponded to the standard Cr_2O_3 , and no any other peak e.g. Cr^{6+} (580.2 eV) is detected, which further demonstrate the substitution of Cr^{3+} cations with Sr^{2+} in the Cr-doped $\text{SrTiO}_3/\text{TiO}_2$ heterostructures^{23,24}. In addition, as shown in Fig. S7, the variation of the Sr 3d peak position is much larger than that of Ti 2p, implying that doping Cr cations into SrTiO_3 mainly affects the Sr sites instead of the Ti sites, which is in good agreement with the above deduced conclusions.

Moreover, the photoconversion efficiencies of as-prepared heterostructured Cr-doped $\text{SrTiO}_3/\text{TiO}_2$ nanotube arrays have been explored. As shown in Fig. 4A, when all these samples were illuminated with continuous monochromatic light, an intense photocurrent peak at around 370 nm can be clearly observed (Fig. 4B), the peak edges of the photocurrent response shifted first to right and then to left by increasing of the hydrothermal time. At the reaction time of 1 h, the largest bathochromic shift can be observed and the peak edge of the photocurrent approximated to 500 nm. More specifically, according to the absorption spectra (Fig. 3A), the absorption peak edges red-shifted continuously, the peak edges of the photocurrent should also shift toward right correspondingly. However, the actual measurements did not really confirm this prediction. It is well known that the photoelectrochemical performances are influenced by not only the excitation and separation of electron-hole pairs but also the transport of charge carriers, and the final photocurrent density were determined by the synergistic function of these two factors. Therefore, although the absorption spectra red shifted continuously with the increase of hydrothermal times, the photoconversion efficiency decreased unexpectedly due to the severe destruction of the TiO_2 tubular structures (Fig. S2), which go against the electron transport. Moreover, it is generally believed that the recombination of

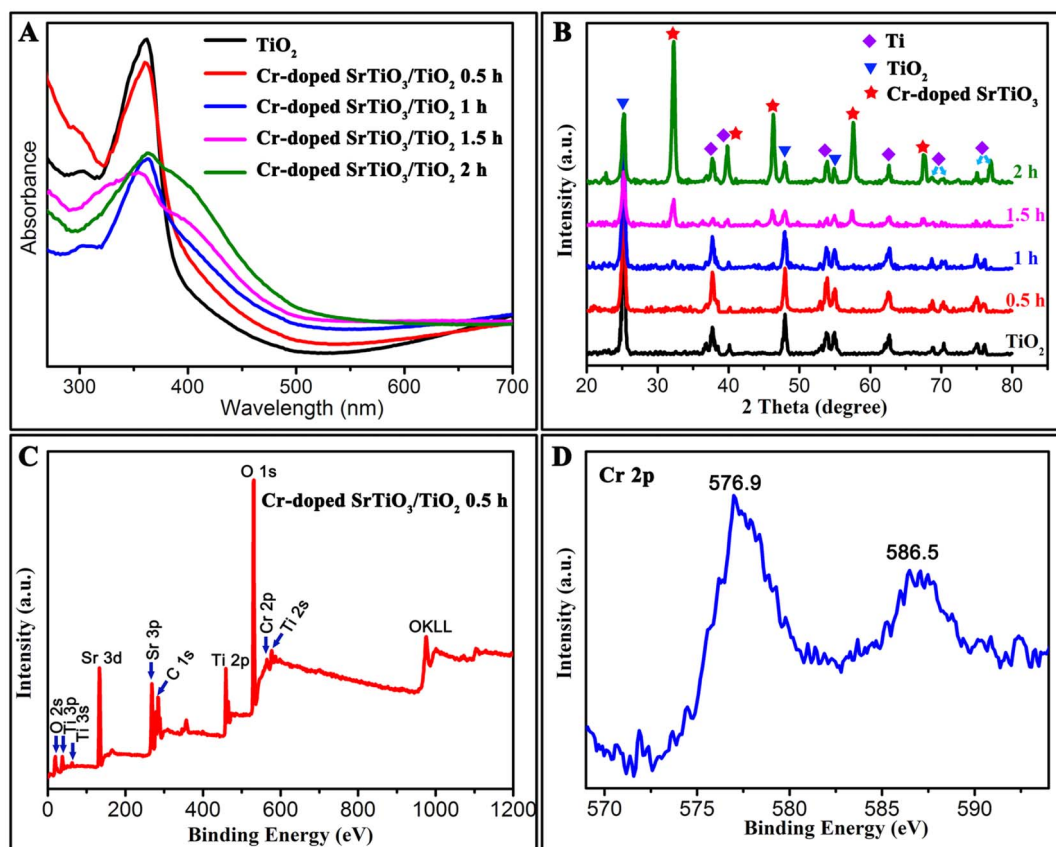


Figure 3 | (A) UV-Vis diffusive absorption spectra and (B) XRD patterns of TiO_2 nanotube arrays and heterostructured Cr-doped $\text{SrTiO}_3/\text{TiO}_2$ nanotube arrays. (C) XPS spectra and (D) high-resolution Cr 2p XPS spectra of the 0.5 h treated sample.

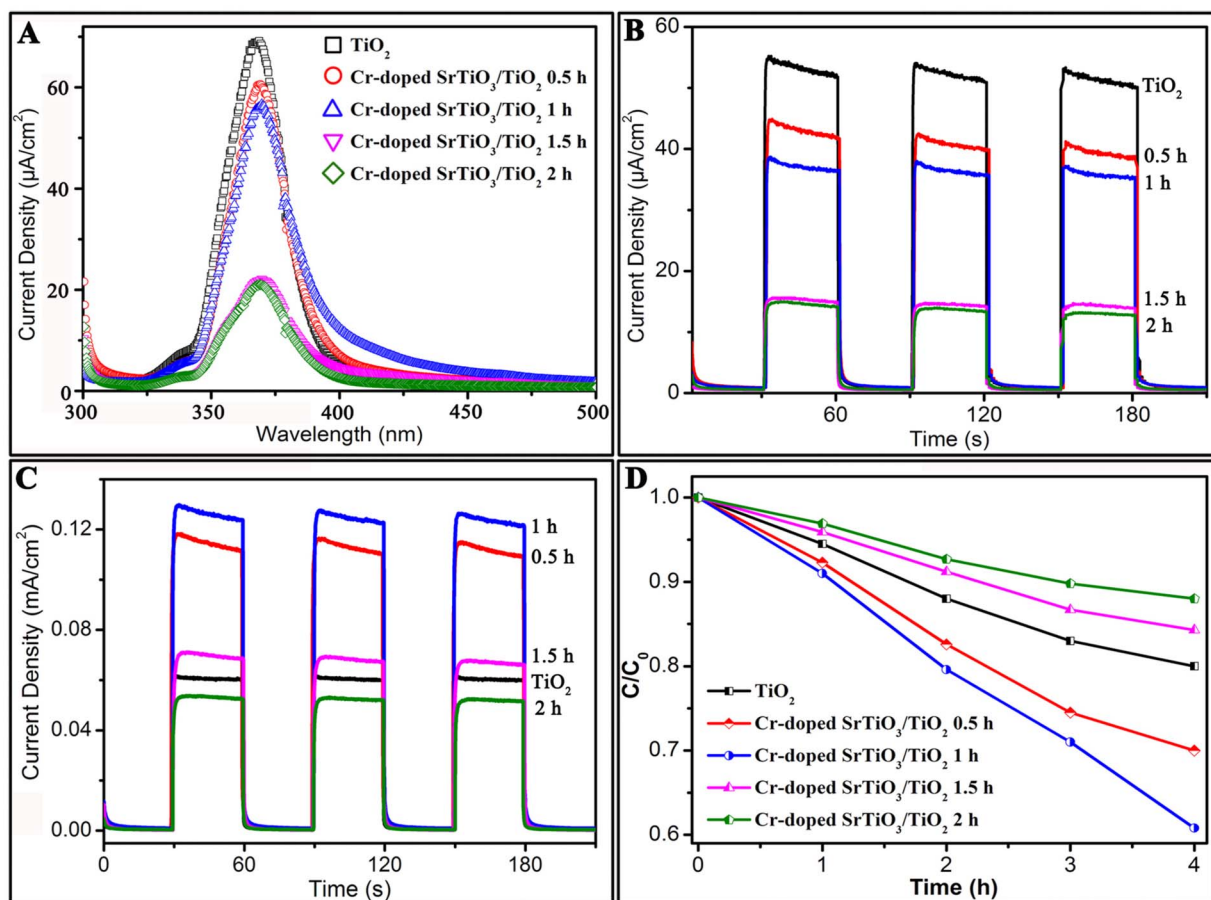


Figure 4 | Photoelectric conversion performances of Cr-doped SrTiO₃/TiO₂ nanotube array under (A) continuous monochromatic light, (B) 372 nm monochromatic light and (C) visible light irradiation (λ > 420 nm) and (D) Photocatalytic performances for RhB degradation under visible light irradiation (λ > 420 nm).

photoinduced electron-hole pairs would become more severe at increased doping level because of the formation of more defects, which might be another unfavourable factor for the photoconversion.

As shown in Fig. 4C, when the samples were illuminated under visible light, except for the 2 h treated sample, all the Cr-doped SrTiO₃/TiO₂ nanocomposites exhibited higher photocurrent properties than pure TiO₂ nanotube arrays, and the photocurrent of the 1 h treated sample was twice as high as that of pure TiO₂. Moreover, it can be found that the photocurrent of the Cr-doped SrTiO₃/TiO₂ heterostructure nanotube arrays increased along with the hydrothermal reaction times before 1 h. With further increasing treatment time over 1 h, the photocurrent density decreased. Hence, it can be deduced that the degree of coverage of Cr-doped SrTiO₃ over TiO₂ plays a crucial role in determining their photoelectrochemical properties. On the other hand, the photocatalytic behaviours of the as-prepared heterostructured Cr-doped SrTiO₃/TiO₂ nanotube arrays for the degradation of RhB under visible light irradiation have also been investigated and shown in Fig. 4D. It can be seen that the photocatalytic activities of these Cr-doped SrTiO₃/TiO₂ nanocomposites were in generally accordance with their photoelectrochemical performances except the 1.5 h treated one, and the 1 h treated sample still exhibited the highest photocatalytic property.

Discussion

According to the discussion above, it can be concluded that the Cr-doped SrTiO₃ greatly determine the photoelectrochemical and photocatalytic performances of TiO₂ nanotube arrays. Figure 5 illustrates the possible mechanism concerning the improved charge separation of Cr-doped SrTiO₃/TiO₂ heterostructure nanotube arrays

under visible light irradiation. The valence band top of SrTiO₃ is lifted up by the occupied Cr³⁺ level, while the original conduction band bottom determined by Ti 3d orbital is hardly affected²². Under visible light illumination, the photogenerated electrons of Cr-doped SrTiO₃ are photoexcited from the valence band (Cr 3d) to the conduction band (Ti 3d) and then transferred to the conduction band of TiO₂ because the conduction band of SrTiO₃ is 200 mV more negative than that of TiO₂^{13–18}. As a result, direct coupling of Cr-doped SrTiO₃ to TiO₂ can not only enhance the visible light response but also reduce the recombination of charge carriers at the surface of the heterostructure. Moreover, the vectorial electron transfer through aligned TiO₂ nanotube arrays facilitates charge transport and thus enhances the photoconversion efficiency²⁵. Therefore, when TiO₂ nanotube arrays were covered by thin and uniform Cr-doped SrTiO₃ nuclei through short duration hydrothermal treatment (≤ 1 h), the photoconversion efficiency of the heterostructured Cr-doped SrTiO₃/TiO₂ nanotube arrays was enhanced greatly. However, further increasing hydrothermal time, TiO₂ nanotubes arrays would be coated with high thick and densely packed Cr-doped SrTiO₃ nuclei, which could severely destroy the tubular structure and hinder the excellent electron transport ability of TiO₂ nanotube arrays. The inner TiO₂, being lost in its transformation to Cr-doped SrTiO₃, would scarcely make any contribution to the transfer and transport of electrons, and the photocurrent response of the nanocomposites decreased accordingly. The photoelectrochemical performances of hetero-semiconductors are the synergistic results of the excitation of electrons, separation of electron-hole pairs and the transport of charge carriers. In the case of photocatalysis, because the photocatalytic activities are significantly influenced by the separation

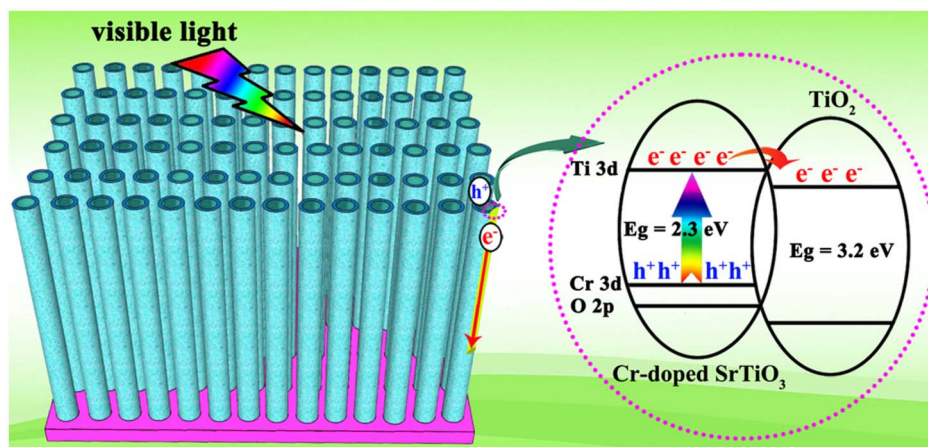


Figure 5 | Schematic illustration for the charge separation of (A) Cr-doped SrTiO₃/TiO₂ heterostructure nanotube arrays.

of electron-hole pairs rather than the transport of charge carriers, the decrease of the photocatalytic performances of these heterostructures may be due to the unfavorable electron transfer and the rapid recombination of electron-hole pairs with the increase of Cr-doped SrTiO₃ and the diminish of TiO₂.

In summary, we have demonstrated a facile and efficient process for fabricating heterostructured Cr-doped SrTiO₃/TiO₂ nanotube arrays. Moreover, their performance studies indicated that the Cr-doped SrTiO₃/TiO₂ heterostructure nanotube arrays exhibited higher photoelectrochemical and photocatalytic activities than pure TiO₂ nanotube arrays and SrTiO₃/TiO₂ heterostructure nanotube arrays without Cr doping under visible light irradiation. The possible function of chromium element for enhancing their both photoelectrochemical and photocatalytic behaviours has also been discussed detailedly. It is considered that this strategy may also be generally adapted for preparing other metal doped TiO₂ based heterostructures, which may have promising applications in both photoelectric conversion and photocatalysis.

Methods

Chemicals. Titanium foil (0.1 mm, > 99.9%) was purchased from Baoji Titanium Industry Co., Ltd. Acetone, ethanol, nitric acid, hydrofluoric acid, ammonium fluoride, chromic nitrate and ethylene glycol were obtained from Sinopharm Chemical Reagent Co., Ltd. Strontium hydroxide octahydrate (Sr(OH)₂·8H₂O) was purchased from Alfa. Deionized water (18 MΩ, Molecular) was used for all solution preparations.

Sample preparation. Firstly, titanium foil (purity > 99.9%) with 0.1 mm thickness was cut into strips with 1 cm width and 3 cm length. Before anodization, these strips were degreased by sonication in acetone, ethanol and deionized water in turn for about 20 min, and then polished in the mixture of hydrofluoric acid and nitric acid at the ratio of 1 : 1. The titanium strip was placed in a plastic beaker equipped with a platinum slice negative electrode and continuous current power supply. Ammonium fluoride (0.3 wt %) and deionized water (3 wt %) diluted in ethylene glycol (200 ml) was used as electrolyte. A constant potential of 60 V was supplied between the two electrodes for 30 min. Subsequently, the strip was washed with ethylene glycol and sonicated for 1 min, and then washed with deionized water and sonicated for 10 s to remove the surface deposit. At last, these strips were annealed at 500°C for 1 h with a programming rate of 3°C/min for both warming up and cooling down. Well aligned crystalline anatase TiO₂ nanotube arrays with the average length of 3.8 μm were obtained.

The annealed TiO₂ nanotube arrays obtained in the above step were used as both substrate and reactant for the fabrication of Cr-doped SrTiO₃/TiO₂ nanocomposites. In general, 5 mol % Cr cations doped SrTiO₃/TiO₂ heterostructure nanotube arrays are fabricated as follows. Firstly, 0.25 g strontium hydroxide (Sr(OH)₂·8H₂O) and 0.02 g chromic nitrate (Cr(NO₃)₃·9H₂O) was added to 40 ml deionized water with vigorous stirring. Next, the solution was transferred into a Teflon-lined stainless steel autoclave which contains a piece of TiO₂ nanotube array strip. The autoclave was kept at 150°C in an oven for 0.5, 1, 1.5, and 2 h, respectively. After the hydrothermal reaction, the strip was drawn from the autoclave and washed with deionized water, and then dried at 60°C for 1 h. Heterostructured Cr-doped SrTiO₃/TiO₂ nanotube arrays have been synthesized.

Materials characterization. The morphology and size of the as-prepared products were characterized by using a field-emission scanning electron microscope (JSM-6701F, JEOL) operated at an accelerating voltage of 5 kV. The X-ray diffraction spectra (XRD) measurements were performed on a PANalytical X'Pert PRO instrument using Cu Kα radiation (40 kV). The XRD patterns were recorded from 20° to 80° with a scanning rate of 0.067°/s. UV-visible diffuse reflectance spectra were taken on a UV-2550 (Shimadzu) spectrometer by using BaSO₄ as the reference. The element composition was detected by X-ray photoelectron spectroscopy (XPS, Kratos Axis Ultra DLD). HAADF imaging and STEM-EDX mapping techniques were carried out using an FEI Tecnai TF20 microscope operated at 200 kV.

Photoelectrochemical and photocatalytic measurements. The TiO₂ nanotube arrays and Cr-doped SrTiO₃/TiO₂ nanocomposites were used as photoanodes in the 3-arm photoelectrochemical cell. Photocurrent response under visible light irradiation were recorded with a CHI-660D potentiostat in a sandwich-type configuration using Pt slice as counter electrode, a saturated calomel electrode (SCE) as the reference, and 0.1 M Na₂SO₄ solution as electrolyte. A 300 W xenon arc lamp equipped with an ultraviolet cutoff filter (HSX-F300, Beijing NBeT Technology Co., Ltd) calibrated to 100 mW/cm², which was measured with a radiometer (CEL-NP2000, Beijing Au-light Co., Ltd), served as a light source. The photocurrent response under monochromatic light measurements was performed using a 300 W Xe lamp and a monochromator (71SWS, Beijing 7-Star Optical Instruments Co., Ltd) at 0.6 V vs SCE. All experiments were carried out under ambient condition.

In all catalytic activity of experiments, 100 ml rhodamine B (RhB) solution with an initial concentration of 0.25 × 10⁻⁵ g/L in the presence of a strip of Cr-doped SrTiO₃/TiO₂ nanotube arrays (1 cm × 3 cm) was filled in a photoreactor, which was then irradiated with a 300 W Xe arc lamp equipped with an ultraviolet cutoff filter. The degradation of organic dyes was monitored by UV/Vis spectroscopy (UV-2550PC, Shimadzu). At given intervals of illumination, the samples of the reaction solution were taken out and analyzed. Decrease in the concentration of dye solution was measured at λ = 554.

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Acknowledgments

This work was supported by the “Hundred Talents Program” of the Chinese Academy of Science and National Natural Science Foundation of China (21273255, 21303232).

Author contributions

Z.B.J. and Y.P.B. developed initial concept. T.C., X.J.Y. and T.W. designed experiments. Z.B.J. and T.C. performed experiments and analyzed data. Z.B.J. wrote the manuscript. G.X.L., J.H.Y. and Y.P.B. supervised the study. All authors discussed the results and commented on the manuscript.

Additional information

Supplementary information accompanies this paper at <http://www.nature.com/scientificreports>

Competing financial interests: The authors declare no competing financial interests.

How to cite this article: Jiao, Z. *et al.* Visible-light-driven photoelectrochemical and photocatalytic performances of Cr-doped SrTiO₃/TiO₂ heterostructured nanotube arrays. *Sci. Rep.* **3**, 2720; DOI:10.1038/srep02720 (2013).



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